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8. [Amended] A pharmaceutical tablet composition comprising:

(a) an effective amount of ibuprofen wherein the weight of the ibuprofen is provided in a range, of the total weight of the tablet composition, of up to about 50%;

(b) an effective amount of hydrocodone;

(c) colloidal silicon dioxide provided in a range, by total weight of the tablet composition, of about 1.5% to about 2%;

(d) microcrystalline cellulose provided in a range, of the total weight of the tablet composition, of about 15% to about 25%;

(e) a disintegrant selected from the group consisting of croscarmellose sodium and crospovidone wherein the weight of the disintegrant is provided in a range, of the total weight of the tablet composition, of about 6 to about 8%;

(f) a binder consisting of an alkylhydroxy methylcellulose wherein the weight of the binder is provided in a range, of the total weight of the tablet composition, of about 3 % to about 4 %;

(g) corn starch wherein the weight of the corn starch is provided in a range, of the total weight of the tablet composition, of about 11 to about 17 %; and

(h) a lubricant wherein the weight of the lubricant is provided in an amount less than 1% by weight of the total weight of the tablet.